From: Rate, Debra
To: Laws, Meredith

Subject: RE: Letter from Dr. Steven Bradbury Concerning Neonicotinoid Registrations

Date: Tuesday, July 30, 2013 1:26:50 PM

Lois agreed that the granular product is not subject to this, but that the SC product would be. She doesn't believe putting a sticker on the product with new labeling would be a problem.

From: Laws, Meredith

Sent: Tuesday, July 30, 2013 9:50 AM

To: Rate, Debra

Subject: FW: Letter from Dr. Steven Bradbury Concerning Neonicotinoid Registrations

I think we can tell him that it was decided that his product is not going to be subject to this – but can you check with Lois?

From: Akers, Rodney [mailto:Rodney.Akers@arysta.com]

Sent: Monday, July 29, 2013 5:07 PM

To: Laws, Meredith

Subject: RE: Letter from Dr. Steven Bradbury Concerning Neonicotinoid Registrations

Hello Meredith,

I was able to postpone the production of our granular product until Sept but our SC product is in production now with bottles and labels ready to go. Can we sticker product in our control with new language? Any ideas of how much language revision EPA is considering? Sorry for the questions, I just have several marketing people upset with me as you might imagine. Thank you for the heads up.

All the best, Rodney Regulatory Affairs Lead



Arysta LifeScience North America 15401 Weston Parkway, Suite 150 Cary, NC 27513

Tel: 919-678-4922 Mobile: 919-946-8509 Fax: 919-678-2183

E-mail: rodney.akers@arysta.com (Please note new email)

From: Laws, Meredith [mailto:Laws.Meredith@epa.gov]

Sent: Monday, July 29, 2013 12:08 PM

To: Akers, Rodney

Subject: RE: Letter from Dr. Steven Bradbury Concerning Neonicotinoid Registrations

Hi Rodney:

An email is fine for submitting information to us. We will be sending the labeling out probably during the week of August 12 so if you can hold off printing until then, that would be best.

Thank you for providing this information, we appreciate it.

Meredith

From: Akers, Rodney [mailto:Rodney.Akers@arysta.com]

Sent: Thursday, July 25, 2013 6:11 PM

To: Laws, Meredith

Subject: RE: Letter from Dr. Steven Bradbury Concerning Neonicotinoid Registrations

Dear Ms. Laws,

I hope this email is OK to respond for the information requested in Dr. Bradbury's letter for production and label printing. We are currently in production now of our products containing clothianidin and plan to formulate again in April and August of 2014. We could print new (revised)labels for the 2014 production if needed. Please let me know if this email is OK or should I submit a letter to you.

Thank you.

Best regards, Rodney Akers Regulatory Affairs Lead



Arysta LifeScience North America 15401 Weston Parkway, Suite 150 Cary, NC 27513

Tel: 919-678-4922 Mobile: 919-946-8509 Fax: 919-678-2183

E-mail: rodney.akers@arysta.com (Please note new email)

From: Laws, Meredith [mailto:Laws.Meredith@epa.gov]

Sent: Monday, July 22, 2013 1:01 PM

Subject: Letter from Dr. Steven Bradbury Concerning Neonicotinoid Registrations

Dear Registrants & Agents:

Please read the attached letter from Dr. Steven Bradbury, Director of EPA's Office of Pesticide Programs. The letter is pertaining to your products containing the nitroguanidine neonicotinoids.

We are asking that you acknowledge receipt of this email within 48 hours.

Thank you,

Meredith Laws
Chief, Insecticide-Rodenticide Branch
Registration Division
Office of Pesticide Programs
(703) 308-7038
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